Metrocare 500 mg tablets for dogs and cats

Authorised

- Metronidazole

Product identification

Medicine name:
Metrocare 500 mg tablets for dogs and cats

Active substance:
Metronidazole

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Metronidazole
500.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01XD01
QP51AA01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:
500 tablets - 50 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.
50 tablets - 5 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.
60 tablets - 6 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.
40 tablets - 4 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.
80 tablets - 8 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.
70 tablets - 7 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.
20 tablets - 2 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.
100 tablets - 10 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.
30 tablets - 3 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.
250 tablets - 25 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.
90 tablets - 9 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.

10 tablets - 1 PVC-Aluminium-Oriented polyamide -Aluminium blister pack in a cardboard box.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ecuphar

Marketing authorisation date:
13/09/2019

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10491/012/002

Date of authorisation status change:
13/09/2019

Reference member state:
Netherlands

Procedure number:
NL/V/0239/002

Concerned member states:
Austria Belgium Czechia Denmark Estonia Finland France Germany Hungary Ireland Luxembourg Norway Poland Portugal Romania Slovakia Spain Sweden United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents